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NEWS 1		Web Page for STN Seminar Schedule - N. America
NEWS 2	APR 04	STN AnaVist, Version 1, to be discontinued
NEWS 3	APR 15	WPIDS, WPINDEX, and WPIX enhanced with new predefined hit display formats
NEWS 4	APR 28	EMBASE Controlled Term thesaurus enhanced
NEWS 5	APR 28	IMSRESEARCH reloaded with enhancements
NEWS 6	MAY 30	INPAFAMDB now available on STN for patent family searching
NEWS 7	MAY 30	DGENE, PCTGEN, and USGENE enhanced with new homology sequence search option
NEWS 8	JUN 06	EPFULL enhanced with 260,000 English abstracts
NEWS 9	JUN 06	KOREAPAT updated with 41,000 documents
NEWS 10	JUN 13	USPATFULL and USPAT2 updated with 11-character patent numbers for U.S. applications
NEWS 11	JUN 19	CAS REGISTRY includes selected substances from web-based collections
NEWS 12	JUN 25	CA/CAplus and USPAT databases updated with IPC reclassification data
NEWS 13	JUN 30	AEROSPACE enhanced with more than 1 million U.S. patent records
NEWS 14	JUN 30	EMBASE, EMBAL, and LEMBASE updated with additional options to display authors and affiliated organizations
NEWS 15	JUN 30	STN on the Web enhanced with new STN AnaVist Assistant and BLAST plug-in
NEWS 16	JUN 30	STN AnaVist enhanced with database content from EPFULL
NEWS 17	JUL 28	CA/CAplus patent coverage enhanced
NEWS 18	JUL 28	EPFULL enhanced with additional legal status information from the epoline Register
NEWS 19	JUL 28	IFICDB, IFIPAT, and IFIUDB reloaded with enhancements
NEWS 20	JUL 28	STN Viewer performance improved
NEWS 21	AUG 01	INPADOCDB and INPAFAMDB coverage enhanced
NEWS 22	AUG 13	CA/CAplus enhanced with printed Chemical Abstracts page images from 1967-1998
NEWS 23	AUG 15	CAOLD to be discontinued on December 31, 2008
NEWS 24	AUG 15	CAplus currency for Korean patents enhanced
NEWS 25	AUG 25	CA/CAplus, CASREACT, and IFI and USPAT databases enhanced for more flexible patent number searching
NEWS 26	AUG 27	CAS definition of basic patents expanded to ensure comprehensive access to substance and sequence information

NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,

AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

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NEWS LOGIN Welcome Banner and News Items
NEWS IPC8 For general information regarding STN implementation of IPC 8

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* * * * * * * * * * * * * * * * STN Columbus * * * * * * * * * * * * * * * *

FILE 'HOME' ENTERED AT 09:42:14 ON 15 SEP 2008

=> fil reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

FILE 'REGISTRY' ENTERED AT 09:42:31 ON 15 SEP 2008
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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 14 SEP 2008 HIGHEST RN 1049627-95-3
DICTIONARY FILE UPDATES: 14 SEP 2008 HIGHEST RN 1049627-95-3

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TSCA INFORMATION NOW CURRENT THROUGH July 5, 2008.

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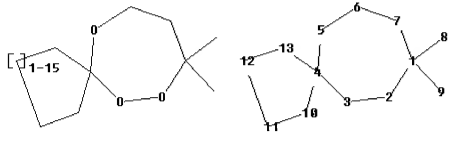
REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

=>
Uploading C:\Program Files\STNEXP\Queries\10564554.str



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chain nodes :
8  9
ring nodes :
1  2  3  4  5  6  7  10  11  12  13
chain bonds :
1-8  1-9
ring bonds :
1-2  1-7  2-3  3-4  4-5  4-10  4-13  5-6  6-7  10-11  11-12  12-13
exact/norm bonds :
1-2  1-7  2-3  3-4  4-5  4-10  4-13  5-6  6-7  10-11  11-12  12-13
exact bonds :
1-8  1-9


Match level :
1:Atom  2:Atom  3:Atom  4:Atom  5:Atom  6:Atom  7:Atom  8:CLASS  9:CLASS  10:Atom
11:Atom  12:Atom  13:Atom



L1      STRUCTURE UPLOADED

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L1 HAS NO ANSWERS
L1                STR
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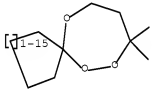


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Structure attributes must be viewed using STN Express query preparation.


=> s sss sam l1
SAMPLE SEARCH INITIATED 09:42:53 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED -       24 TO ITERATE

100.0% PROCESSED       24 ITERATIONS                                3 ANSWERS
SEARCH TIME: 00.00.01

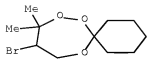
FULL FILE PROJECTIONS:  ONLINE  **COMPLETE**
                        BATCH   **COMPLETE**
PROJECTED ITERATIONS:          187 TO      773
PROJECTED ANSWERS:               3 TO      163

L2            3 SEA SSS SAM L1

=> d scan

L2   3 ANSWERS   REGISTRY  COPYRIGHT 2008 ACS on STN
```

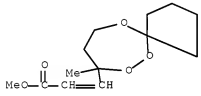
IN 7,8,12-Trioxaspiro[5.6]dodecane, 10-bromo-9,9-dimethyl-
MF C11 H19 Br O3



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):3

L2 3 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN 2-Propenoic acid, 3-(8-methyl-6,7,11-trioxaspiro[4.6]undec-8-yl)-, methyl ester
MF C13 H20 O5



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 3 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
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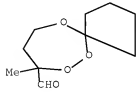
MF C10 H16 O4



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED

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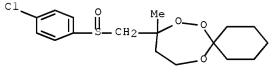
FULL SEARCH INITIATED 09:45:02 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 399 TO ITERATE

100.0% PROCESSED 399 ITERATIONS 34 ANSWERS
SEARCH TIME: 00.00.01

L3 34 SEA SSS FUL L1

=> d scan

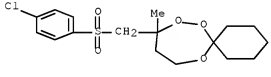
L3 34 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN 7,8,12-Trioxaspiro[5.6]dodecane, 9-[[(4-chlorophenyl)sulfinyl]methyl]-9-methyl-
MF C17 H23 Cl O4 S



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

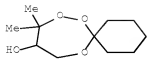
HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):5

L3 34 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN 7,8,12-Trioxaspiro[5.6]dodecane, 9-[[(4-chlorophenyl)sulfonyl]methyl]-9-methyl-
MF C17 H23 Cl O5 S



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

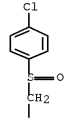
L3 34 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN 7,8,12-Trioxaspiro[5.6]dodecan-10-ol, 9,9-dimethyl-
MF C11 H20 O4



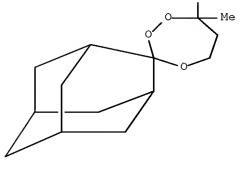
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L3 34 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN Spiro[tricyclo[3.3.1.13,7]decane-2,3'-[1,2,4]trioxepane],
7'-[[(4-chlorophenyl)sulfinyl]methyl]-7'-methyl-
MF C21 H27 Cl O4 S

PAGE 1-A



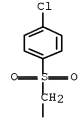
PAGE 2-A



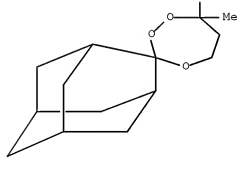
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L3 34 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN Spiro[tricyclo[3.3.1.13,7]decane-2,3'-[1,2,4]trioxepane],
7'-[[(4-chlorophenyl)sulfonyl]methyl]-7'-methyl-
MF C21 H27 Cl O5 S

PAGE 1-A

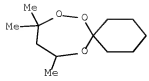


PAGE 2-A



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L3 34 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN 7,8,12-Trioxaspiro[5.6]dodecane, 9,9,11-trimethyl-
MF C12 H22 O3

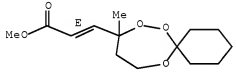


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L3 34 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
IN 2-Propenoic acid, 3-(9-methyl-7,8,12-trioxaspiro[5.6]dodec-9-yl)-, methyl ester, (2E)-
MF C14 H22 O5

Double bond geometry as shown.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):end

=> save temp l3 meij10564554/a
ANSWER SET L3 HAS BEEN SAVED AS 'MEIJ10564554/A'

=> fil caplu
COST IN U.S. DOLLARS

	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	180.66	180.87

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FILE COVERS 1907 - 15 Sep 2008 VOL 149 ISS 12
FILE LAST UPDATED: 14 Sep 2008 (20080914/ED)

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```
=> s l3 (L) ract/rl
             9 L3
       3157313 RACT/RL
L4           2 L3 (L) RACT/RL

=> s l3
L5           9 L3

=> s l3/rrt
'RRT' IS NOT A VALID CROSSOVER QUALIFIER FOR L3
Answer sets created in a different file may be field qualified with a
limited set of qualifiers.  Enter HELP CROSSOVER at an arrow prompt
(=>) for specific information.

=> fil casreact
COST IN U.S. DOLLARS                          SINCE FILE      TOTAL
                                                 ENTRY    SESSION
FULL ESTIMATED COST                               3.08     183.95

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FILE CONTENT:1840 - 13 Sep 2008 VOL 149 ISS 12

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```
*************************************************************
*                                                           *
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*                                                           *
*************************************************************
```

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This file contains CAS Registry Numbers for easy and accurate substance identification.

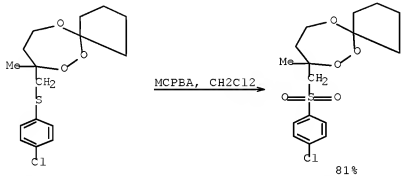
=> s l3/rrt
L6 3 L3/RRT

=> d sca

L6 3 ANSWERS CASREACT COPYRIGHT 2008 ACS on STN

TI Synthesis of 1,2,4-trioxepanes via application of thiol-olefin Co-oxygenation methodology

RX(4) OF 52



81%

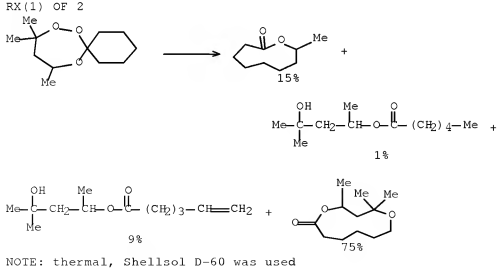
NOTE: product depends on equiv. of m-CPBA, safety

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L6 3 ANSWERS CASREACT COPYRIGHT 2008 ACS on STN

TI 1,2,4,-Trioxepanes as precursors for lactones

RX(1) OF 2



NOTE: thermal, Shellsol D-60 was used

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

=> fil caplu
COST IN U.S. DOLLARS

	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	31.03	214.98

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FILE LAST UPDATED: 14 Sep 2008 (20080914/ED)

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http://www.cas.org/legal/infopolicy.html

=> s l4 or l5 or l6
3 L6
L7 9 L4 OR L5 OR L6

=> d ibib abs hitst 1-9
'HITST' IS NOT A VALID FORMAT FOR FILE 'CAPLUS'

The following are valid formats:

ABS --------- GI and AB
ALL --------- BIB, AB, IND, RE
APPS -------- AI, PRAI
BIB --------- AN, plus Bibliographic Data and PI table (default)
CAN --------- List of CA abstract numbers without answer numbers
CBIB -------- AN, plus Compressed Bibliographic Data
CLASS ------- IPC, NCL, ECLA, FTERM
DALL -------- ALL, delimited (end of each field identified)
DMAX -------- MAX, delimited for post-processing
FAM --------- AN, PI and PRAI in table, plus Patent Family data
FBIB -------- AN, BIB, plus Patent FAM
IND --------- Indexing data
IPC --------- International Patent Classifications
MAX --------- ALL, plus Patent FAM, RE
PATS -------- PI, SO

```
SAM --------- CC, SX, TI, ST, IT
SCAN -------- CC, SX, TI, ST, IT (random display, no answer numbers;
              SCAN must be entered on the same line as the DISPLAY,
              e.g., D SCAN or DISPLAY SCAN)
STD --------- BIB, CLASS

IABS -------- ABS, indented with text labels
IALL -------- ALL, indented with text labels
IBIB -------- BIB, indented with text labels
IMAX -------- MAX, indented with text labels
ISTD -------- STD, indented with text labels

OBIB -------- AN, plus Bibliographic Data (original)
OIBIB ------- OBIB, indented with text labels

SBIB -------- BIB, no citations
SIBIB ------- IBIB, no citations

HIT --------- Fields containing hit terms
HITIND ------ IC, ICA, ICI, NCL, CC and index field (ST and IT)
              containing hit terms
HITRN ------- HIT RN and its text modification
HITSTR ------ HIT RN, its text modification, its CA index name, and
              its structure diagram
HITSEQ ------ HIT RN, its text modification, its CA index name, its
              structure diagram, plus NTE and SEQ fields
FHITSTR ----- First HIT RN, its text modification, its CA index name, and
              its structure diagram
FHITSEQ ----- First HIT RN, its text modification, its CA index name, its
              structure diagram, plus NTE and SEQ fields
KWIC -------- Hit term plus 20 words on either side
OCC --------- Number of occurrence of hit term and field in which it occurs

To display a particular field or fields, enter the display field
codes.  For a list of the display field codes, enter HELP DFIELDS at
an arrow prompt (=>).  Examples of formats include: TI; TI,AU; BIB,ST;
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information will be displayed in the same order as the format
specification.

All of the formats (except for SAM, SCAN, HIT, HITIND, HITRN, HITSTR,
FHITSTR, HITSEQ, FHITSEQ, KWIC, and OCC) may be used with DISPLAY ACC
to view a specified Accession Number.
ENTER DISPLAY FORMAT (BIB):end

=> d ibib abs hitstr l7 1-9

L7   ANSWER 1 OF 9  CAPLUS  COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:         2007:1421759  CAPLUS  Full-text
DOCUMENT NUMBER:          148:55109
TITLE:                    Preparation of dual molecules containing a peroxide
                          derivative, particularly 1,2,4-
                          trioxane/trioxepane/trioxecane, linked to a quinoline
                          moiety as antimalarial agents
INVENTOR(S):              Cosledan, Frederic; Meunier, Bernard; Pellet, Alain
PATENT ASSIGNEE(S):       Sanofi Aventis, Fr.; Palumed; Centre National de la
                          Recherche Scientifique (C.N.R.S.)
SOURCE:                   Fr. Demande, 53pp.
                          CODEN: FRXXBL
DOCUMENT TYPE:            Patent
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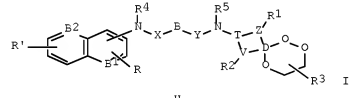
LANGUAGE: French
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
FR 2902100	A1	20071214	FR 2006-5235	20060613
WO 2007144487	A2	20071221	WO 2007-FR946	20070608
WO 2007144487	A3	20080207		

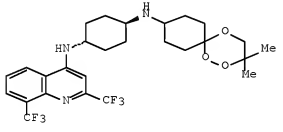
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA

PRIORITY APPLN. INFO.: FR 2006-5235 A 20060613
OTHER SOURCE(S): CASREACT 148:55109; MARPAT 148:55109
GI



I



II

AB Trioxanes of formula I and related derivs. [R, R' = independently H, halo, OH, cycloalkyl, NO2, NH2, etc.; B1 = N and B2 = CH: or B1 = CH: and B2 = N; R4 = H, alkyl, cycloalkyl; B = (un)substituted cycloalkyl or 2 cycloalkyl groups containing C3-6 rings connected by a bond, CH2, or CH2-CH2; X = (CH2)m; Y = (CH2)n; m, n = independently 0-2; R5 = H, (cyclo)alkyl, CO-(cyclo)alkyl, COO-(cyclo)alkyl; Z, V = independently alkylene with Z+V+T+D = cycloalkyl, polycyclyl; one of Z or V = a bond between the T and D carbon atoms with the proviso that Z and V cannot simultaneously be a bond; R1, R2 = independently H, a functional group capable of increasing solubility in water; D = junction carbon; each R3 = independently H, halo, OH, CF3, aryl, cycloalkyl, bi- or tricyclyl, etc.; or R3CCR3 = (un)substituted C5-C6 cycloalkyl; or R3CR3 C3-C7 cycloalkyl or C4-C18 bi- or tricyclyl], and their acid addition salts, hydrates and solvates, and their stereoisomers were prepared as antimalarial agents. Thus, amination of 4-chloro-2,8-bis(trifluoromethyl)quinoline with

trans-1,4- diaminocyclohexane, and reductive amination of 3,3-dimethyl-1,2,5-trioxaspiro[5.5]undecan-9-one (preparation given) with the primary amine intermediate gave spiro-1,2,4-trioxane II. I were tested in vitro for antimalarial activity against 2 strains of P. falciparum, i.e. FcB1-Columbia and FcM29-CAmeroon and displayed IC50 values < 1 μM. I displayed a high metabolic stability in the human hepatic microsomes.

IT 960114-71-0P

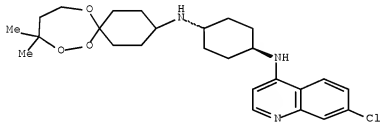
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of quinolinyl trioxanes, trioxepanes and trioxecanes as antimalarial agents)

RN 960114-71-0 CAPLUS

CN 1,4-Cyclohexanediamine, N1-(7-chloro-4-quinolinyl)-N4-(9,9-dimethyl-7,8,12-trioxaspiro[5.6]dodec-3-yl)-, trans- (CA INDEX NAME)

Relative stereochemistry.



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 2 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2006:1190098 CAPLUS Full-text

DOCUMENT NUMBER: 146:142841

TITLE: Synthesis of 1,2,4-trioxepanes via application of thiol-olefin Co-oxygenation methodology

AUTHOR(S): Amewu, Richard; Stachulski, Andrew V.; Berry, Neil G.; Ward, Stephen A.; Davies, Jill; Labat, Gael; Rossignol, Jean-Francois; O'Neill, Paul M.

CORPORATE SOURCE: Department of Chemistry, University of Liverpool, Liverpool, L69 3BX, UK

SOURCE: Bioorganic & Medicinal Chemistry Letters (2006), 16(23), 6124-6130
CODEN: BMCLE8; ISSN: 0960-894X

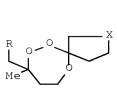
PUBLISHER: Elsevier Ltd.

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 146:142841

GI



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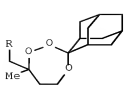
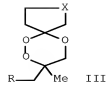


II



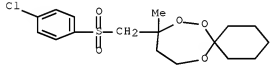
III

AB Thiol-olefin co-oxygenation (TOCO) of substituted allylic alcs. generated β-hydroxy peroxides that can be condensed in situ with various ketones, to afford a series of functionalized 1,2,4-trioxepanes in good yields. Manipulation of the phenylsulfenyl group in I [R = SC6H4-4-Cl, X = (CH2)n, n = 1, 2] and II (R = SC6H4-4-Cl) allowed for convenient modification to the spiro-trioxepane substituents. Surprisingly, and in contrast to the 1,2,4-trioxanes examined, the 1,2,4-trioxepanes were inactive as antimalarials up to 1000 nM, an observation based the inherent stability of these systems to ferrous mediated degradation FMO calcns. clearly show that the σ* orbital of the peroxide moiety of 1,2,4-trioxane derivs. III [R = SC6H4-4-Cl, SO2C6H4-4-Cl, X = (CH2)2] are lower in energy and more accessible to attack by Fe(II) compared to their trioxepane analogs I [R = SC6H4-4-Cl, SO2C6H4-4-Cl, X = (CH2)2].

IT 869661-34-7P 869661-35-8P 918901-86-7P
918901-92-5P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)
(synthesis and antimalarial activity of 1,2,4-trioxepane artemisinin analogs via application of thiol-olefin co-oxygenation methodol.)

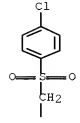
RN 869661-34-7 CAPLUS

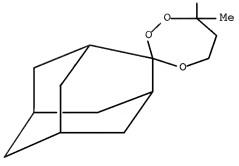
CN 7,8,12-Trioxaspiro[5.6]dodecane, 9-[[(4-chlorophenyl)sulfonyl]methyl]-9-methyl- (CA INDEX NAME)



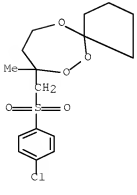
RN 869661-35-8 CAPLUS

CN Spiro[tricyclo[3.3.1.13,7]decane-2,3'-[1,2,4]trioxepane], 7'-[[(4-chlorophenyl)sulfonyl]methyl]-7'-methyl- (CA INDEX NAME)



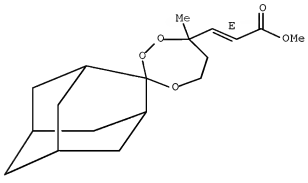


RN 918901-86-7 CAPLUS
CN 6,7,11-Trioxaspiro[4.6]undecane, 8-[[(4-chlorophenyl)sulfonyl]methyl]-8-methyl- (CA INDEX NAME)



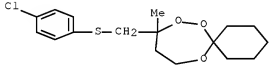
RN 918901-92-5 CAPLUS
CN 2-Propenoic acid, 3-(7'-methylspiro[tricyclo[3.3.1.13,7]decane-2,3'-[1,2,4]trioxepan]-7'-yl)-, methyl ester, (2E)- (CA INDEX NAME)

Double bond geometry as shown.

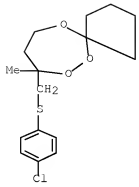


IT 869661-31-4P 869661-32-5P 869661-33-6P
918901-87-8P 918901-88-9P 918901-94-7P
918901-95-8P 918902-01-9P 918902-02-0P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(synthesis and antimalarial activity of 1,2,4-trioxepane artemisinin analogs via application of thiol-olefin co-oxygenation methodol.)

RN 869661-31-4 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane, 9-[[(4-chlorophenyl)thio]methyl]-9-methyl- (CA INDEX NAME)



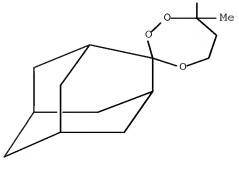
RN 869661-32-5 CAPLUS
CN 6,7,11-Trioxaspiro[4.6]undecane, 8-[[(4-chlorophenyl)thio]methyl]-8-methyl- (CA INDEX NAME)



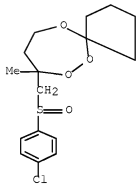
RN 869661-33-6 CAPLUS
CN Spiro[tricyclo[3.3.1.13,7]decane-2,3'-[1,2,4]trioxepane], 7'-[[(4-chlorophenyl)thio]methyl]-7'-methyl- (CA INDEX NAME)

PAGE 1-A

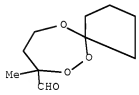
PAGE 2-A



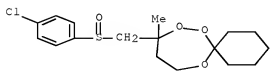
RN 918901-87-8 CAPLUS
CN 6,7,11-Trioxaspiro[4.6]undecane, 8-[[(4-chlorophenyl)sulfinyl]methyl]-8-methyl- (CA INDEX NAME)



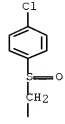
RN 918901-88-9 CAPLUS
CN 6,7,11-Trioxaspiro[4.6]undecane-8-carboxaldehyde, 8-methyl- (CA INDEX NAME)

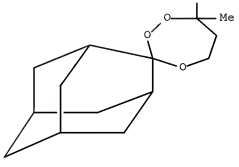


RN 918901-94-7 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane, 9-[[(4-chlorophenyl)sulfinyl]methyl]-9-methyl- (CA INDEX NAME)

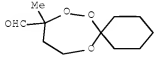


RN 918901-95-8 CAPLUS
CN Spiro[tricyclo[3.3.1.13,7]decane-2,3'-[1,2,4]trioxepane], 7'-[[(4-chlorophenyl)sulfinyl]methyl]-7'-methyl- (CA INDEX NAME)

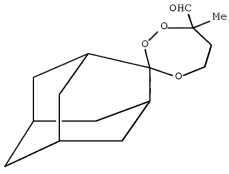




RN 918902-01-9 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane-9-carboxaldehyde, 9-methyl- (CA INDEX NAME)

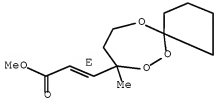


RN 918902-02-0 CAPLUS
CN Spiro[tricyclo[3.3.1.13,7]decane-2,3'-[1,2,4]trioxepane]-7'-carboxaldehyde, 7'-methyl- (CA INDEX NAME)



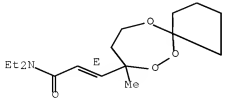
IT 918901-89-0P 918901-90-3P 918901-96-9P
918901-97-0P 918901-98-1P
RL: SPN (Synthetic preparation); PREP (Preparation)
(synthesis and antimalarial activity of 1,2,4-trioxepane artemisinin analogs via application of thiol-olefin co-oxygenation methodol.)
RN 918901-89-0 CAPLUS
CN 2-Propenoic acid, 3-(8-methyl-6,7,11-trioxaspiro[4.6]undec-8-yl)-, methyl ester, (2E)- (CA INDEX NAME)

Double bond geometry as shown.



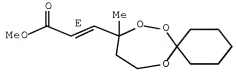
RN 918901-90-3 CAPLUS
CN 2-Propenamide, N,N-diethyl-3-(8-methyl-6,7,11-trioxaspiro[4.6]undec-8-yl)-, (2E)- (CA INDEX NAME)

Double bond geometry as shown.



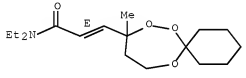
RN 918901-96-9 CAPLUS
CN 2-Propenoic acid, 3-(9-methyl-7,8,12-trioxaspiro[5.6]dodec-9-yl)-, methyl ester, (2E)- (CA INDEX NAME)

Double bond geometry as shown.



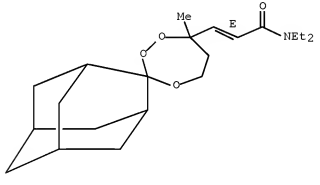
RN 918901-97-0 CAPLUS
CN 2-Propenamide, N,N-diethyl-3-(9-methyl-7,8,12-trioxaspiro[5.6]dodec-9-yl)-, (2E)- (CA INDEX NAME)

Double bond geometry as shown.



RN 918901-98-1 CAPLUS
CN 2-Propenamide, N,N-diethyl-3-(7'-methylspiro[tricyclo[3.3.1.13,7]decane-2,3'-[1,2,4]trioxepan]-7'-yl)-, (2E)- (CA INDEX NAME)

Double bond geometry as shown.



REFERENCE COUNT: 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

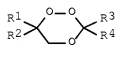
L7 ANSWER 3 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:1223637 CAPLUS Full-text
DOCUMENT NUMBER: 143:477995
TITLE: 1,2,4-Trioxanes and 1,2,4-trioxepanes useful as antimalarial and anticancer agents, and their pharmaceutical compositions, use, and preparation via the thiol-olefin co-oxygenation (TOCO) reaction

INVENTOR(S): O'Neill, Paul M.; Amewu, Richard; Mukhtar, Amira; Ward, Stephen A.
PATENT ASSIGNEE(S): UK
SOURCE: U.S. Pat. Appl. Publ., 19 pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 20050256184	A1	20051117	US 2005-103076	20050411
WO 2006016903	A2	20060216	WO 2005-US12236	20050412
WO 2006016903	A3	20060526		
WO 2006016903	A8	20060720		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRIORITY APPLN. INFO.: US 2004-561589P P 20040412
US 2005-103076 A 20050411
OTHER SOURCE(S): CASREACT 143:477995; MARPAT 143:477995
GI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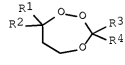
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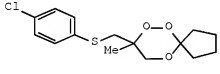
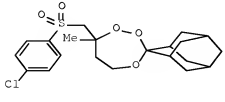

III

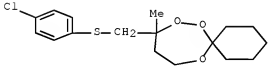

IV

AB Novel substituted 1,2,4-trioxanes and 1,2,4-trioxepanes useful as anti-malarial and/or (no data) anticancer agents, and an improved method for their preparation, are disclosed. The method preferably involves a thiol-olefin co-

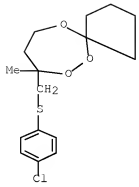
oxygenation (TOCO) reaction between an aromatic thiol, an allylic or homoallylic alc., O2, and a ketone. The trioxanes and trioxepanes are represented by structures I and II, resp. [R1 = CH2OH, CHO, alkenyl, arylsulfonylmethyl, methylsulfinyl, methylpiperazinyl; R2 = aryl, alkyl; R3 = alkyl, cycloalkyl; R4 = alkyl, cycloalkyl; or R3R4 = atoms to form cycloalkyl ring; including enantiomers, salts, and hydrates]. For instance, 2-methyl-2-propen-1-ol and AIBN in MeCN were irradiated with UV light under pure O2 with simultaneous addition of 4-ClC6H4SH, followed by stirring to completion. The intermediate hydroperoxy alc. was treated in situ with cyclopentanone and tosic acid catalyst at -10° to give after workup 40% invention compound III. Oxidation of III with mCPBA gave the corresponding sulfone, also an invention compound, in 96% yield. Analogous reactions starting from homoallylic alcs. gave trioxepanes such as IV. Claimed synthetic methods involve the above-illustrated thiol-olefin co-oxygenation (TOCO) reaction, as well as auxiliary reactions, including the Pummerer reaction, Wittig reaction, condensations, nucleophilic substitutions, and reductive aminations. All of the compds. displayed moderate antimalarial activity, and one or two compds. displayed potent activity, approaching that of artemisinin, against chloroquine-resistant Kl Plasmodium falciparum. X-ray crystal structures were obtained for both III and the dechloro trioxane analog of IV.

IT 869661-31-4P, 9-[[(4-Chlorophenyl)sulfanyl]methyl]-9-methyl-7,8,12-trioxaspiro[5.6]dodecane 869661-32-5P, 8-[[(4-Chlorophenyl)sulfanyl]methyl]-8-methyl-6,7,11-trioxaspiro[4.6]undecane 869661-33-6P 869661-34-7P, 9-[[(4-Chlorophenyl)sulfonyl]methyl]-9-methyl-7,8,12-trioxaspiro[5.6]dodecane 869661-35-8P 869661-36-9P, 9-[2-(Methoxycarbonyl)vinyl]-9-methyl-7,8,12-trioxaspiro[5.6]dodecane 869661-37-0P, 8-[2-(Methoxycarbonyl)vinyl]-8-methyl-6,7,11-trioxaspiro[4.6]undecane 869661-38-1P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of trioxanes and trioxepanes as antimalarial and anticancer agents)

RN 869661-31-4 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane, 9-[[(4-chlorophenyl)thio]methyl]-9-methyl- (CA INDEX NAME)



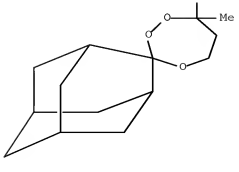
RN 869661-32-5 CAPLUS
CN 6,7,11-Trioxaspiro[4.6]undecane, 8-[[(4-chlorophenyl)thio]methyl]-8-methyl- (CA INDEX NAME)



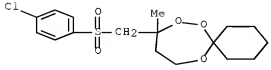
RN 869661-33-6 CAPLUS
CN Spiro[tricyclo[3.3.1.13,7]decane-2,3'-[1,2,4]trioxepane], 7'-[[(4-chlorophenyl)thio]methyl]-7'-methyl- (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

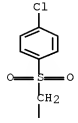


RN 869661-34-7 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane, 9-[[(4-chlorophenyl)sulfonyl]methyl]-9-methyl- (CA INDEX NAME)

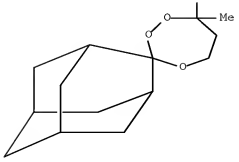


RN 869661-35-8 CAPLUS
CN Spiro[tricyclo[3.3.1.13,7]decane-2,3'-[1,2,4]trioxepane], 7'-[[(4-chlorophenyl)sulfonyl]methyl]-7'-methyl- (CA INDEX NAME)

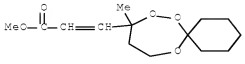
PAGE 1-A



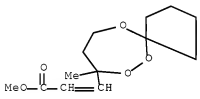
PAGE 2-A



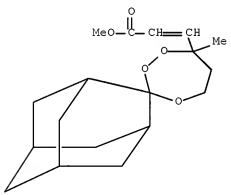
RN 869661-36-9 CAPLUS
CN 2-Propenoic acid, 3-(9-methyl-7,8,12-trioxaspiro[5.6]dodec-9-yl)-, methyl ester (CA INDEX NAME)



RN 869661-37-0 CAPLUS
CN 2-Propenoic acid, 3-(8-methyl-6,7,11-trioxaspiro[4.6]undec-8-yl)-, methyl ester (CA INDEX NAME)



RN 869661-38-1 CAPLUS
CN 2-Propenoic acid, 3-(7'-methylspiro[tricyclo[3.3.1.13,7]decane-2,3'-[1,2,4]trioxepan]-7'-yl)-, methyl ester (CA INDEX NAME)

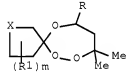


L7 ANSWER 4 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:141049 CAPLUS Full-text
DOCUMENT NUMBER: 142:240473
TITLE: 1,2,4,-Trioxepanes as precursors for lactones
INVENTOR(S): Meijer, John; Van den Berg, Rolf Hendrik

PATENT ASSIGNEE(S): Akzo Nobel N. V., Neth.
SOURCE: PCT Int. Appl., 18 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005014569	A1	20050217	WO 2004-EP7839	20040712
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2004263256	A1	20050217	AU 2004-263256	20040712
EP 1646618	A1	20060419	EP 2004-763237	20040712
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK				
CN 1823053	A	20060823	CN 2004-80020641	20040712
IN 2006CN00182	A	20070817	IN 2006-CN182	20060113
US 20060167281	A1	20060727	US 2006-564554	20060202
PRIORITY APPLN. INFO.:			EP 2003-77238 A	20030717
			US 2003-499415P P	20030902
			WO 2004-EP7839 W	20040712

OTHER SOURCE(S): CASREACT 142:240473; MARPAT 142:240473
GI



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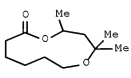


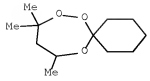
II

AB A novel process was disclosed for the preparation of cyclic ether lactones by decomposition of corresponding 1,2,4-trioxepanes, such as I [R = H, Me; R1 = H, carboxy, etc.; X = (CH2)n, n = 1-14]. Thus, macrocyclic lactone ether II was prepared as the major product of heating 1,2,4-trioxepane I [R = Me, R1 = H, X = (CH2)2, m = 0] in a reactor charged with Shellsol D-60.

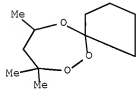
IT 215877-52-4 844665-36-7 844665-37-8
844665-38-9 844665-39-0
RL: RCT (Reactant); PACT (Reactant or reagent)
(process for the preparation of macrocyclic ether lactones via thermal decomposition of spirocyclic 1,2,4,-trioxepanes)

RN 215877-52-4 CAPLUS

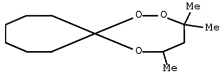
CN 7,8,12-Trioxaspiro[5.6]dodecane, 9,9,11-trimethyl- (CA INDEX NAME)



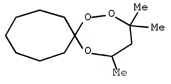
RN 844665-36-7 CAPLUS
CN 6,7,11-Trioxaspiro[4.6]undecane, 8,8,10-trimethyl- (CA INDEX NAME)



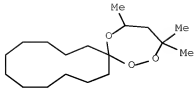
RN 844665-37-8 CAPLUS
CN 1,2,6-Trioxaspiro[6.6]tridecane, 3,3,5-trimethyl- (CA INDEX NAME)



RN 844665-38-9 CAPLUS
CN 1,2,6-Trioxaspiro[6.7]tetradecane, 3,3,5-trimethyl- (CA INDEX NAME)



RN 844665-39-0 CAPLUS
CN 1,2,6-Trioxaspiro[6.11]octadecane, 3,3,5-trimethyl- (CA INDEX NAME)



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

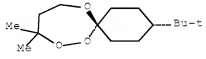
L7 ANSWER 5 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2004:737076 CAPLUS Full-text
DOCUMENT NUMBER: 141:379904
TITLE: 1,2,4-Trioxepanes: Redox-cleavable protection for carbonyl groups
AUTHOR(S): Ahmed, Aqeel; Dussault, Patrick H.
CORPORATE SOURCE: Department of Chemistry, University of Nebraska-Lincoln, Lincoln, NE, 68588-0304, USA
SOURCE: Organic Letters (2004), 6(20), 3609-3611
CODEN: ORLEF7; ISSN: 1523-7060
PUBLISHER: American Chemical Society
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 141:379904

AB 1,2,4-Trioxepanes, readily prepared and easily handled derivs. of aldehydes and ketones, are stable to a variety of synthetic conditions and yet easily deblocked with Zn/HOAc or Mg/MeOH to regenerate the parent carbonyl. Trioxepanes may provide an alternative to 1,3-dithianes for acid-stable protection of carbonyl groups.

IT 784191-69-1P 784191-70-4P
RL: SPN (Synthetic preparation); PREP (Preparation)
(Co-catalyzed cyclization of aldehydes and ketones with (triethylsilylperoxy)methylbutanol to give trioxepanes useful as acid-stable, redox-cleavable carbonyl protecting groups)

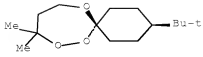
RN 784191-69-1 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane, 3-(1,1-dimethylethyl)-9,9-dimethyl-, (3α,6α)- (CA INDEX NAME)

Relative stereochemistry.



RN 784191-70-4 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane, 3-(1,1-dimethylethyl)-9,9-dimethyl-, (3α,6β)- (CA INDEX NAME)

Relative stereochemistry.

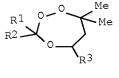


REFERENCE COUNT: 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 6 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2002:142767 CAPLUS Full-text
DOCUMENT NUMBER: 136:200613
TITLE: Use of trioxepanes in the process to modify (co)polymers
INVENTOR(S): Gerritsen, Rene; Hogt, Andreas Herman; Meijer, John
PATENT ASSIGNEE(S): Akzo Nobel N.V., Neth.
SOURCE: PCT Int. Appl., 25 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002014383	A1	20020221	WO 2001-EP9266	20010808
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2002012134	A	20020225	AU 2002-12134	20010808
US 20020040108	A1	20020404	US 2001-930402	20010815
PRIORITY APPLN. INFO.:			US 2000-225314P P	20000815
			EP 2000-203888 A	20001108
			WO 2001-EP9266 W	20010808

OTHER SOURCE(S): MARPAT 136:200613
GI



I

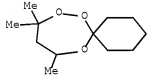
AB Invention relates to a polymer modification process wherein the rheol. of at least one (co)polymers is modified by contacting the (co)polymer with at least one decomposing peroxide of the formula (I), wherein R1-3 = independently

selected from substituted or unsubstituted hydrocarbyl groups. The modification process can be useful to obtain a modified resin or to enhance the flame retardancy of (expanded) styrenic resins. Thus, peroxide I, wherein R1-3 = Me dissolved in dichloromethane (5% weight solution) and Borealis HC 00A1B1 (PP) were mixed in amount such that 0.05% weight active oxygen was introduced, 0.1% weight based on PP Irganox 1010 stabilizer was added, and extruded to give a modified resin with MFR (ASTM D 1238) 82 g/10 min.

IT 215877-52-4
RL: MOA (Modifier or additive use); USES (Uses)
(peroxide; use of trioxepanes in process to modify (co)polymers)

RN 215877-52-4 CAPLUS

CN 7,8,12-Trioxaspiro[5.6]dodecane, 9,9,11-trimethyl- (CA INDEX NAME)



REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 7 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1998:745028 CAPLUS Full-text

DOCUMENT NUMBER: 130:4473

TITLE: Crosslinking thermoplastic polymers using cyclic peroxide and a promoter

INVENTOR(S): Bock, Lawrence A.; Lewis, Roger N.

PATENT ASSIGNEE(S): Witco Corp., USA

SOURCE: PCT Int. Appl., 21 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9850354	A1	19981112	WO 1998-US8907	19980501
W: KR				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
US 5856412	A	19990105	US 1997-850334	19970502
EP 909269	A1	19990421	EP 1998-920129	19980501
EP 909269	B1	20050209		
R: BE, DE, ES, FR, GB, IT, NL, SE				
JP 11106521	A	19990420	JP 1998-123268	19980506
JP 4098879	B2	20080611		
KR 2000022442	A	20000425	KR 1998-710875	19981230
PRIORITY APPLN. INFO.:			US 1997-850334 A	19970502
			WO 1998-US8907 W	19980501

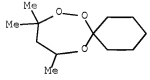
OTHER SOURCE(S): MARPAT 130:4473

AB A crosslinking system includes a 1,2,4-trioxacycloheptane and a crosslink-promoting polyfunctional ethylenically unsatd. compound Preferred polymers for crosslinking, include ethylene homopolymers and copolymers. Thus, HDPE was cured in the presence of triallyl cyanurate and 1.1% 9,9,11-trimethyl-7,8,12-

trioxaspiro[5.6]dodecane (preparation given) at 200°, 30 min showing crosslink degree 44.1% (based on weight of product).

IT 215877-52-4P
RL: CAT (Catalyst use); IMF (Industrial manufacture); PREP (Preparation); USES (Uses)
(crosslinking catalyst; crosslinking thermoplastic polymers using cyclic peroxide and a unsatd. compound promoter)

RN 215877-52-4 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane, 9,9,11-trimethyl- (CA INDEX NAME)

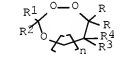


REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

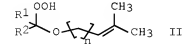
L7 ANSWER 8 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER:	1997:790896 CAPLUS Full-text
DOCUMENT NUMBER:	128:75374
ORIGINAL REFERENCE NO.:	128:14743a,14746a
TITLE:	New methods for the synthesis of oxy-functionalized 1,2,4-trioxanes and 1,2,4-trioxepanes from unsaturated hydroperoxy acetals
AUTHOR(S):	Ushigoe, Yoshihiro; Masuyama, Araki; Nojima, Masatomo; Mccullough, Kevin J.
CORPORATE SOURCE:	Department of Materials Chemistry, Faculty of Engineering, Osaka University, Suita, 565, Japan
SOURCE:	Tetrahedron Letters (1997), 38(50), 8753-8756 CODEN: TELEAY; ISSN: 0040-4039
PUBLISHER:	Elsevier Science Ltd.
DOCUMENT TYPE:	Journal
LANGUAGE:	English
OTHER SOURCE(S):	CASREACT 128:75374

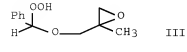
GI



I



II



III

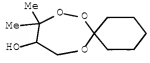
AB Title compds. I (R = Me, H, CH2OH, C(CH2)2OH; R1 = Ph; R2 = H; R1R2 = (CH2)5; R3 = H, OH; R4 =H, Me; n = 0, 1, 2) were prepared from II by autoxidn. or from III, prepared from II, by acid-catalyzed cyclization.

IT 200639-92-5P

RL: SPN (Synthetic preparation); PREP (Preparation)
(synthesis of oxy-functionalized 1,2,4-trioxanes and 1,2,4-trioxepanes from unsatd. hydroperoxy acetals)

RN 200639-92-5 CAPLUS

CN 7,8,12-Trioxaspiro[5.6]dodecan-10-ol, 9,9-dimethyl- (CA INDEX NAME)



REFERENCE COUNT: 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 9 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1997:81259 CAPLUS Full-text

DOCUMENT NUMBER: 126:199546

ORIGINAL REFERENCE NO.: 126:38579a

TITLE: Synthesis of 1,2,4-trioxanes and 1,2,4-trioxepanes by N-halogenosuccinimide-mediated cyclizations of unsaturated hydroperoxyacetals

AUTHOR(S): Ushigoe, Yoshihiro; Kano, Yoshihiro; Nojima, Masatomo

CORPORATE SOURCE: Fac. Eng., Osaka Univ., Osaka, 565, Japan

SOURCE: Journal of the Chemical Society, Perkin Transactions 1: Organic and Bio-Organic Chemistry (1997), (1), 5-10
CODEN: JCPRB4; ISSN: 0300-922X

PUBLISHER: Royal Society of Chemistry

DOCUMENT TYPE: Journal

LANGUAGE: English

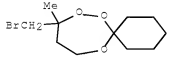
OTHER SOURCE(S): CASREACT 126:199546

GI

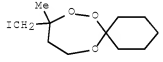
AB Ozonolyses of vinyl ethers R1R2C:CHOMe [R1 = Ph, C7H15, R2 = H; R1R2 = (CH2)5] in CH2Cl2 in the presence of allylic and homoallylic alcs. R42C:CR3CH2OH (R3 = Me, H, R4 = H, Me) and H2C:CMeCH2CH2OH give in each case the corresponding unsatd. hydroperoxy acetals R42C:CR3CH2OCR1R2OOH and H2C:CMeCH2CH2OCR1R2OOH, derived from capture of the carbonyl oxides by the unsatd. alcs. N-Halosuccinimide-mediated cyclizations of the hydroperoxides give the corresponding 1,2,4-trioxanes, e.g., I and/or 1,2,4-trioxepanes, e.g., II (X = Br, iodo), depending on the structure of the hydroperoxides and the identity of the N-halosuccinimides.

IT 187884-37-3P 187884-38-4P 187884-46-4P
187884-48-6P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of trioxanes and trioxepanes by cyclization of unsatd. hydroperoxyacetals)

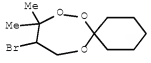
RN 187884-37-3 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane, 9-(bromomethyl)-9-methyl- (CA INDEX NAME)



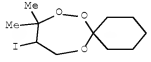
RN 187884-38-4 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane, 9-(iodomethyl)-9-methyl- (CA INDEX NAME)



RN 187884-46-4 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane, 10-bromo-9,9-dimethyl- (CA INDEX NAME)



RN 187884-48-6 CAPLUS
CN 7,8,12-Trioxaspiro[5.6]dodecane, 10-iodo-9,9-dimethyl- (CA INDEX NAME)



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